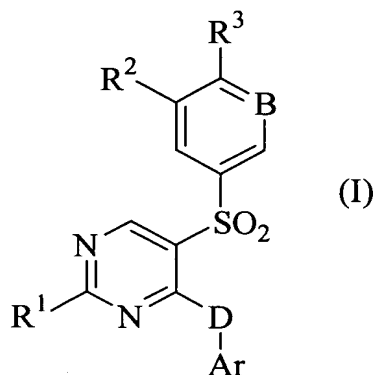


What is claimed is:

1. A compound of Formula (I)



5

or pharmaceutically acceptable salt or solvate thereof,

wherein

10

B is CH or N;

D is CH<sub>2</sub> or NH;

15

R<sup>1</sup> is selected from the group consisting of H, -CN, C<sub>1-4</sub> alkyl, C<sub>3-7</sub> cycloalkyl, C<sub>2-4</sub> alkenyl, C<sub>2-4</sub> alkynyl, C<sub>1-4</sub> alkoxy and N(C<sub>1-4</sub> alkyl)<sub>2</sub> optionally and independently substituted with 1 to 3 substituents selected from the group consisting of -CN, hydroxy, halo, C<sub>1-4</sub> haloalkyl and C<sub>1-4</sub> alkoxy;

20

R<sup>2</sup> is selected from the group consisting of H, halo, -CN, hydroxy, C<sub>1-6</sub> alkyl, C<sub>2-6</sub> alkenyl, C<sub>2-6</sub> alkynyl, C<sub>3-7</sub> cycloalkyl, C<sub>1-6</sub> alkoxy, C<sub>1-6</sub> haloalkyl, -NR<sup>4</sup>R<sup>6</sup>, -C<sub>1-6</sub>alkylNR<sup>4</sup>R<sup>6</sup>, -C<sub>1-6</sub>alkylOR<sup>6</sup>, CO<sub>2</sub>R<sup>6</sup>, O<sub>2</sub>CR<sup>6</sup>, COR<sup>6</sup>, CON<sup>4</sup>R<sup>6</sup>, NR<sup>4</sup>CO<sub>2</sub>R<sup>6</sup>, NR<sup>4</sup>SO<sub>2</sub>R<sup>6</sup>, NR<sup>4</sup>COR<sup>6</sup>, OCONR<sup>4</sup>R<sup>6</sup> and NR<sup>4</sup>CONR<sup>5</sup>R<sup>6</sup>;

25

optionally and independently substituted with 1  
to 3 substituents selected from the group  
consisting of -CN, hydroxy, halo, C<sub>1-4</sub>  
haloalkyl, C<sub>1-4</sub> alkoxy, CO<sub>2</sub>C<sub>1-4</sub> alkyl or  
phenyl; or

R<sup>2</sup> is morpholinyl, thiomorpholinyl,  
piperadiny, piperaziny, phenyl, pyridyl,  
pyrimidinyl, triazinyl, quinolinyl,  
isoquinolinyl, thienyl, imidazolyl,  
thiazolyl, indolyl, pyrrolyl,  
pyrrolidinyl, dihydroimidazolyl, oxazolyl,  
benzofuranyl, benzothienyl,  
benzothiazolyl, benzoxazolyl, isoxazolyl,  
triazolyl, tetrazolyl and indazolyl,  
independently and optionally substituted  
with 1 to 4 substituents selected from the  
group consisting of H, C<sub>1-6</sub> alkyl, C<sub>1-4</sub>  
alkoxy- C<sub>1-4</sub> alkyl, C<sub>3-6</sub> cycloalkyl, -OR<sup>4</sup>,  
halo, C<sub>1-4</sub> haloalkyl, -CN, SH, -S(O)<sub>2</sub>R<sup>5</sup>,  
-COR<sup>4</sup>, -CO<sub>2</sub>R<sup>4</sup>, -OC(O)R<sup>5</sup>, -N(COR<sup>4</sup>)<sub>2</sub>, -NR<sup>4</sup>R<sup>7</sup>  
and -CONR<sup>4</sup>R<sup>7</sup>, -NR<sup>4</sup>COR<sup>5</sup>, NR<sup>4</sup>SO<sub>2</sub>R<sup>5</sup>, NR<sup>4</sup>CONR<sup>5</sup>R<sup>7</sup>  
or NR<sup>4</sup>CO<sub>2</sub>R<sup>5</sup>;

R<sup>3</sup> is selected from the group consisting of H, halo,  
-CN, hydroxy, C<sub>1-6</sub> alkyl, C<sub>2-6</sub> alkenyl, C<sub>2-6</sub>  
alkynyl, C<sub>3-7</sub> cycloalkyl, C<sub>1-6</sub> alkoxy, C<sub>1-6</sub>  
haloalkyl, -NR<sup>4</sup>R<sup>6</sup>, -C<sub>1-6</sub>alkylNR<sup>4</sup>R<sup>6</sup>, -C<sub>1-6</sub>alkylOR<sup>6</sup>,  
CO<sub>2</sub>R<sup>6</sup>, O<sub>2</sub>CR<sup>6</sup>, COR<sup>6</sup>, CON<sup>4</sup>R<sup>6</sup>, NR<sup>4</sup>CO<sub>2</sub>R<sup>6</sup>, NR<sup>4</sup>SO<sub>2</sub>R<sup>6</sup>,  
NR<sup>4</sup>COR<sup>6</sup>, OCONR<sup>4</sup>R<sup>6</sup>, and NR<sup>4</sup>CONR<sup>5</sup>R<sup>6</sup>;

optionally and independently substituted with 1  
to 3 substituents selected from the group  
consisting of -CN, hydroxy, halo, C<sub>1-4</sub>

haloalkyl, C<sub>1-4</sub> alkoxy, CO<sub>2</sub>C<sub>1-4</sub> alkyl,  
phenyl or naphthyl; or

R<sup>3</sup> is morpholinyl, thiomorpholinyl,  
piperadinyl, piperazinyl, phenyl, pyridyl,  
5 pyrimidinyl, triazinyl, quinolinyl,  
isoquinolinyl, thienyl, imidazolyl,  
thiazolyl, indolyl, pyrrolyl,  
pyrrolidinyl, dihydroimidazolyl, oxazolyl,  
benzofuranyl, benzothienyl,  
10 benzothiazolyl, benzoxazolyl, isoxazolyl,  
triazolyl, tetrazolyl and indazolyl,  
independently and optionally substituted  
with 1 to 4 substituents selected from the  
group consisting of H, C<sub>1-6</sub> alkyl, C<sub>3-6</sub>  
15 cycloalkyl, C<sub>1-4</sub> alkoxy- C<sub>1-4</sub> alkyl, -OR<sup>4</sup>,  
halo, C<sub>1-4</sub> haloalkyl, -CN, SH, -S(O)<sub>2</sub>R<sup>5</sup>,  
-COR<sup>4</sup>, -CO<sub>2</sub>R<sup>4</sup>, -OC(O)R<sup>5</sup>, -N(COR<sup>4</sup>)<sub>2</sub>, -NR<sup>4</sup>R<sup>7</sup>  
and -CONR<sup>4</sup>R<sup>7</sup>, -NR<sup>4</sup>COR<sup>5</sup>, NR<sup>4</sup>SO<sub>2</sub>R<sup>5</sup>, NR<sup>4</sup>CONR<sup>5</sup>R<sup>7</sup>  
or NR<sup>4</sup>CO<sub>2</sub>R<sup>5</sup>;

20 Ar is selected from the group consisting of phenyl,  
indanyl, indenyl, pyridyl, pyrimidinyl,  
triazinyl, furanyl, quinolinyl, isoquinolinyl,  
thienyl, imidazolyl, thiazolyl, indolyl,  
pyrrolyl, pyrrolidinyl, dihydroimidazolyl,  
25 oxazolyl, benzofuranyl, benzothienyl,  
benzothiazolyl, benzoxazolyl, isoxazolyl,  
triazolyl, tetrazolyl, indazolyl, indolinyl,  
benzoxazolin-2-on-yl, benzodioxolanyl and  
benzodioxane, independently and optionally  
30 substituted with 1 to 4 substituents selected  
from the group consisting of H, C<sub>1-6</sub> alkyl, C<sub>3-6</sub>  
cycloalkyl, C<sub>1-4</sub> alkoxy- C<sub>1-4</sub> alkyl, -OR<sup>4</sup>, halo,

C<sub>1-4</sub> haloalkyl, -CN, -NO<sub>2</sub>, SH, -S(O)<sub>2</sub>R<sup>5</sup>, -COR<sup>4</sup>,  
 -CO<sub>2</sub>R<sup>4</sup>, -OC(O)R<sup>5</sup>, -N(COR<sup>4</sup>)<sub>2</sub>, -NR<sup>4</sup>R<sup>7</sup> and -CONR<sup>4</sup>R<sup>7</sup>,  
 -NR<sup>4</sup>COR<sup>5</sup>, NR<sup>4</sup>SO<sub>2</sub>R<sup>5</sup>, NR<sup>4</sup>CONR<sup>5</sup>R<sup>7</sup>, and NR<sup>4</sup>CO<sub>2</sub>R<sup>5</sup>;

5 R<sup>4</sup>, R<sup>5</sup> and R<sup>7</sup> are independently selected from the  
 group consisting of H, C<sub>1-6</sub> alkyl, C<sub>3-6</sub>  
 cycloalkyl, C<sub>3-6</sub> cycloalkyl-C<sub>3-6</sub> alkyl, C<sub>1-2</sub>  
 alkoxy-C<sub>1-4</sub> alkyl and C<sub>1-4</sub> haloalkyl; and

10 R<sup>6</sup> is selected from the group consisting of H, C<sub>1-6</sub>  
 alkyl, C<sub>3-6</sub> cycloalkyl, C<sub>3-6</sub> cycloalkyl-C<sub>1-6</sub>  
 alkyl, C<sub>1-2</sub> alkoxy-C<sub>1-2</sub> alkyl, C<sub>1-4</sub> haloalkyl,  
 phenyl and C<sub>1-6</sub> alkyl-phenyl.

2. A compound according to claim 1 wherein B is CH.

15 3. A compound according to claim 1 wherein B is CH and  
 D is CH<sub>2</sub>.

4. A compound according to claim 1 wherein B is CH and  
 D is NH.

20 5. A compound according to claim 1 wherein R<sup>1</sup> is C<sub>1-4</sub>  
 alkyl.

25 6. A compound according to claim 1 wherein R<sup>2</sup> is H or  
 substituted or unsubstituted C<sub>1-6</sub>alkyl, morpholinyl,  
 piperazinyl or phenyl.

7. A compound according to claim 1 wherein R<sup>3</sup> is H,  
 halo, CN or hydroxy, substituted or unsubstituted C<sub>1-6</sub>  
 alkyl, C<sub>1-6</sub> alkoxy, C<sub>1-6</sub> haloalkyl, -NR<sup>4</sup>R<sup>6</sup> or O<sub>2</sub>CR<sup>6</sup>.

30

8. A compound according to claim 1 wherein R<sup>3</sup> is pyrimidinyl and pyridinyl.
9. A compound according to claim 1 wherein Ar is  
5 phenyl, pyridyl, pyrimidinyl, imidazolyl, thiazolyl, pyrrolidinyl, dihydroimidazolyl optionally substituted with 1 to 4 substituents selected from the group consisting of H, C<sub>1-6</sub> alkyl, -OR<sup>4</sup>, halo, C<sub>1-4</sub> haloalkyl, -CN, -NO<sub>2</sub> or -CO<sub>2</sub>R<sup>4</sup>.
10. A compound according to claim 1 wherein R<sup>4</sup>, R<sup>5</sup> and R<sup>7</sup> are independently H or C<sub>1-6</sub> alkyl.
11. A compound according to claim 1 wherein R<sup>6</sup> is H.
12. A compound according to claim 1 wherein R<sup>1</sup> is C<sub>1-4</sub> alkyl; R<sup>2</sup> is H or substituted or unsubstituted C<sub>1-6</sub>alkyl, morpholinyl, piperazinyl or phenyl; R<sup>3</sup> is H, halo, CN or hydroxy, substituted or unsubstituted C<sub>1-6</sub> alkyl, C<sub>1-6</sub>  
20 alkoxy, C<sub>1-6</sub> haloalkyl, -NR<sup>4</sup>R<sup>6</sup> or O<sub>2</sub>CR<sup>6</sup>; Ar is phenyl, pyridyl, pyrimidinyl, imidazolyl, thiazolyl, pyrrolidinyl, dihydroimidazolyl optionally substituted with 1 to 4 substituents selected from the group consisting of H, C<sub>1-6</sub> alkyl, -OR<sup>4</sup>, halo, C<sub>1-4</sub> haloalkyl,  
25 -CN, -NO<sub>2</sub> or -CO<sub>2</sub>R<sup>4</sup>; R<sup>4</sup>, R<sup>5</sup> and R<sup>7</sup> are independently H or C<sub>1-6</sub> alkyl; and R<sup>6</sup> is H.
13. A compound according to claim 1 wherein B is CH; R<sup>1</sup> is C<sub>1-4</sub> alkyl; R<sup>2</sup> is H or substituted or unsubstituted C<sub>1-6</sub>alkyl, morpholinyl, piperazinyl or phenyl; R<sup>3</sup> is H,  
30 halo, CN or hydroxy, substituted or unsubstituted C<sub>1-6</sub> alkyl, C<sub>1-6</sub> alkoxy, C<sub>1-6</sub> haloalkyl, -NR<sup>4</sup>R<sup>6</sup> or O<sub>2</sub>CR<sup>6</sup>; Ar is phenyl, pyridyl, pyrimidinyl, imidazolyl, thiazolyl,

pyrrolidinyl, dihydroimidazolyl optionally substituted with 1 to 4 substituents selected from the group consisting of H, C<sub>1-6</sub> alkyl, -OR<sup>4</sup>, halo, C<sub>1-4</sub> haloalkyl, -CN, -NO<sub>2</sub> or -CO<sub>2</sub>R<sup>4</sup>; R<sup>4</sup>, R<sup>5</sup> and R<sup>7</sup> are independently H or  
 5 C<sub>1-6</sub> alkyl; and R<sup>6</sup> is H.

14. [5-(4-Methoxybenzenesulfonyl)-2-methylpyrimidin-4-yl]-(2,4,6-trimethylphenyl)-amine; 4-[2-Methyl-4-(2,4,6-trimethylphenylamino)-pyrimidine-5-sulfonyl]-phenol;
- 10 Acetic acid 4-[2-methyl-4-(2,4,6-trimethylphenylamino)-pyrimidine-5-sulfonyl]-phenyl ester; [5-(4-Benzyloxybenzenesulfonyl)-2-methylpyrimidin-4-yl]-(2,4,6-trimethylphenyl)-amine; [5-(4-Benzyloxybenzenesulfonyl)-2-methylpyrimidin-4-yl]-(4-methoxy-2-methylphenyl)-amine;
- 15 [5-(4-Benzyloxybenzenesulfonyl)-2-methylpyrimidin-4-yl]-(6-methoxy-2-methylpyridin-3-yl)-amine; [5-(3-Benzyloxybenzenesulfonyl)-2-methylpyrimidin-4-yl]-(2,4,6-trimethylphenyl)-amine; [5-(3-Benzyloxybenzenesulfonyl)-2-methoxypyrimidin-4-yl]-(2,4,6-trimethylphenyl)-amine;
- 20 5-(3-Benzyloxybenzenesulfonyl)-N<sup>2</sup>,N<sup>2</sup>-dimethyl-N<sup>4</sup>-(2,4,6-trimethylphenyl)-pyrimidine-2,4-diamine; {5-[4-(2-Methoxybenzyloxy)-benzenesulfonyl]-2-methylpyrimidin-4-yl]-(2,4,6-trimethylphenyl)-amine; {5-[4-(3,5-Dimethoxybenzyloxy)-benzenesulfonyl]-2-methylpyrimidin-4-yl]-(2,4,6-trimethylphenyl)-amine; [5-(4-Benzyloxybenzenesulfonyl)-2-methylpyrimidin-4-yl]-(2,4-dimethoxyphenyl)-amine; 5-(4-Methoxyoxybenzenesulfonyl)-2-methyl-4-(2,4,6-trimethylbenzyl)-pyrimidine; 5-(4-Benzyloxybenzenesulfonyl)-2-methyl-4-(2,4,6-trimethylbenzyl)-pyrimidine; [5-(4-Fluorobenzenesulfonyl)-2-methylpyrimidin-4-yl]-(2,4,6-trimethylphenyl)-amine; [2-Methyl-5-(4-morpholin-4-yl-benzenesulfonyl)-pyrimidin-4-yl]-(2,4,6-trimethylphenyl)-
- 25
- 30

- amine; {2-Methyl-5-[4-(4-methylpiperazin-1-yl)-benzenesulfonyl]-pyrimidin-4-yl}-(2,4,6-trimethylphenyl)-amine; [5-(4-Imidazol-1-yl-benzenesulfonyl)-2-methylpyrimidin-4-yl}-(2,4,6-trimethylphenyl)-amine; [2-Methyl-5-(4-pyrrolidin-1-yl-benzenesulfonyl)-pyrimidin-4-yl}-(2,4,6-trimethylphenyl)-amine; [5-(4-Benzylaminobenzenesulfonyl)-2-methylpyrimidin-4-yl}-(2,4,6-trimethylphenyl)-amine; {5-[4-(Benzylmethylamino)-benzenesulfonyl]-2-methylpyrimidin-4-yl}-(2,4,6-trimethylphenyl)-amine; 4-[2-Methyl-4-(2,4,6-trimethylphenylamino)-pyrimidine-5-sulfonyl]-benzonitrile; [2-Methyl-5-(toluene-4-sulfonyl)-pyrimidin-4-yl}-(2,4,6-trimethylphenyl)-amine; [2-Methyl-5-(4-pyrimidin-5-yl-benzenesulfonyl)-pyrimidin-4-yl}-(2,4,6-trimethylphenyl)-amine; [2-Methyl-5-(4-pyrimidin-2-yl-benzenesulfonyl)-pyrimidin-4-yl}-(2,4,6-trimethylphenyl)-amine; [2-Methyl-5-(4-pyridin-4-yl-benzenesulfonyl)-pyrimidin-4-yl}-(2,4,6-trimethylphenyl)-amine; [2-Methyl-5-(4-pyridin-2-yl-benzenesulfonyl)-pyrimidin-4-yl}-(2,4,6-trimethylphenyl)-amine; [2-Methyl-5-(4-pyridin-3-yl-benzenesulfonyl)-pyrimidin-4-yl}-(2,4,6-trimethylphenyl)-amine;
- {5-[4-(4,5-Dihydro-1H-imidazol-2-yl)-benzenesulfonyl]-2-methyl-pyrimidin-4-yl}-(2,4,6-trimethylphenyl)-amine; or {5-[4-(1H-Imidazol-2-yl)-benzenesulfonyl]-2-methyl-pyrimidin-4-yl}-(2,4,6-trimethylphenyl)-amine or pharmaceutically acceptable salts or solvates thereof.
15. A pharmaceutical composition of a compound according to claim 1.

16. A method of treating depression or anxiety comprising a compound of claim 15.